

Effective Viscosity of Confined Hydrocarbons

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We present molecular dynamics friction calculations for confined hydrocarbon films with molecular lengths from 20 to 1400 carbon atoms. We find that the logarithm of the effective viscosity η_{eff} for nanometer-thin films depends linearly on the logarithm of the shear rate: $\log \eta_{\text{eff}} = C - n \log \dot{\gamma}$, where n varies from 1 (solidlike friction) at very low temperatures to 0 (Newtonian liquid) at very high temperatures, following an inverse sigmoidal curve. Only the shortest chain molecules melt, whereas the longer ones only show a softening in the studied temperature interval $0 < T < 900$ K. The results are important for the frictional properties of very thin (nanometer) films and to estimate their thermal durability.

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The frictional and rheological properties of thin confined fluid films are of great importance in a number of engineering and scientific applications, e.g., in the context of lubrication. Thus, recent studies [1] have shown that for sliding contacts lubricated by organic or silicon oil bulk shear thinning can have a great influence on the friction at intermediate sliding velocities (mixed lubrication). When wetting fluids are confined between solid walls at nanometer separation they often acquire solidlike properties and an increasing squeezing force is necessary in order to reduce the film thickness. If the solid walls are smooth, the fluid molecules arrange in layers parallel to the solid walls and the squeezeout occurs in a quantized way by removing one monolayer after another with increasing pressure [2–4]. Sometimes the last one or two monolayers are so strongly bound that they cannot be removed by squeezing alone. Fluid films confined at the nanometer length scale exhibit viscosity enhancement and nonlinear flow properties characteristic of sheared supercooled liquids approaching their glass transition [5,6].

Experiments for a large variety of fluids (including hydrocarbon fluids and silicon oil) [4,7] have shown that the logarithm of the effective viscosity η_{eff} for nanometer-thin films (typically 3 or 4 monolayers and contact pressures of the order of a few MPa) depends linearly on the logarithm (here and elsewhere with 10 as the basis) of the shear rate: $\log \eta_{\text{eff}} = C - n \log \dot{\gamma}$. If η_{eff} and $\dot{\gamma}$ are measured in SI units, for a large variety of fluids (at room temperature) $C \approx 5$ and $n \approx 0.9$. This linear relation has also been established in other experiments [8,9] and in computer simulations [10,11].

We have performed a very extensive set of molecular dynamics simulations to probe the frictional properties of thin layers of confined hydrocarbon molecules (with molecular lengths from 20 to 1400 carbon atoms). Some of these results may also be relevant for polymer-on-polymer

systems, as recent studies [12] have shown that in this case the shear deformations are localized to a band of material about 2.5 nm thick. Our results for n and C agree with the experimental observation at room temperature, but show that when the temperature increases n varies from 1 (solidlike friction) at very low temperatures to 0 (Newtonian liquid) at very high temperatures.

Our model is similar to those described in Refs. [13,14]. We consider a block and a substrate with atomically flat surfaces separated by a polymer slab consisting of hydrocarbons with molecular lengths 20, 100, and 1400 carbon atoms. The solid walls are treated as single layers of “atoms” bound to rigid flat surfaces by springs corresponding to the long-range elastic properties of 50 Å thick solid slabs, similar to our earlier papers (see, e.g., Refs. [15–17]). The simulation box in the x - y dimension is equal to $124.8 \text{ Å} \times 124.8 \text{ Å}$. In the following, periodic boundary conditions are assumed in the xy plane.

Initially, about half of the molecules are adsorbed on the block surface and half on the substrate surface. Two solids with adsorbed polymer slabs were put into contact and when the temperature was equal to the thermostat temperature everywhere we started to move the upper block surface. The temperature was varied from 0 to 900 K to study the effect of temperature (and also melting) on the shear stress. In our simulations, the polymer films are very thin (~ 3 nm), and the solid walls are connected to a thermostat at a short distance from the polymer slab. Under these circumstances we find that frictional heating effects are not important, and the effective temperature in the polymer film is always close to the thermostat temperature.

Linear alkanes C_nH_{2n+2} (with $n = 20, 100, \text{ and } 1400$) were used as “lubricant” in the present calculations. The CH_2/CH_3 beads are treated in the united atom representation [18,19]. The Lennard-Jones potential was used to model the interaction between beads of different chains

$$U(r) = 4\epsilon_0 \left[\left(\frac{r_0}{r} \right)^{12} - \left(\frac{r_0}{r} \right)^6 \right], \quad (1)$$

and the same potential with modified parameters (ϵ_1 , r_1) was used for the interaction of each bead with the substrate and block atoms.

The parameters were $\epsilon_0 = 5.12$ meV for both the interior and the end beads, and $r_0 = 3.905$ Å. For the interactions within the C_nH_{2n+2} molecules we used the standard optimized potentials for liquid simulations model [18,19], including flexible bonds, bond bending, and torsion interaction, which results in bulk properties in good agreement with experimental data far below the boiling point [20]. Atomic mass 14 (for interior CH_2 beads) and 15 (for the CH_3 end groups) were used. Within a C_nH_{2n+2} chain we assume nearest neighbor C atoms are connected via springs with the spring constant k , which was chosen equal to 10 N/m. Note that this value is 1 order of magnitude smaller than the optimized 450 N/m [18], and was chosen such to facilitate a reasonable time step of 1 fs. We used an angle bending interaction of the form $E(\cos\theta)/k_B = (1/2)k_{\text{bend}}(\cos\theta - \cos\theta_0)^2$ with $k_{\text{bend}} = 62\,543$ K and $\theta_0 = 2.0001$ rad. For the dihedral interaction, we used the functional form in term of a cosine Fourier series $E(\phi)/k_B = \sum_{i=0}^3 c_i \cos^i(\phi)$ with parameters $c_0 = 1009.99$ K, $c_1 = 2018.95$ K, $c_2 = 136.37$ K, $c_3 = -3165.30$ K. Internal beads of separation greater than 3 units are treated similarly as beads from different chains. The number of molecules was equal to 1000, 200, and 14 for the $C_{20}H_{42}$, $C_{100}H_{202}$, and $C_{1400}H_{2802}$ systems, respectively. The hydrocarbon films at room temperature consisted of 6 to 8 monolayers of molecules between the solid surfaces. The (nominal) squeezing pressure p_0 was usually 10 MPa.

We have chosen the polymer-wall atom bond to be so strong that no slip occurs at these interfaces. This is the case with $r_1 = 2.92$ Å, $\epsilon_1 = 160$ meV. The lattice spacings of the block and of the substrate are $a = b = 2.6$ Å.

If v is the sliding velocity and d the film thickness, we define the shear rate $\dot{\gamma} = v/d$ and the effective viscosity $\eta_{\text{eff}} = \sigma_f / \dot{\gamma}$, where σ_f is the frictional shear stress. Figure 1 shows the logarithm of the effective viscosity as a function of the temperature [21] for the (a) $C_{20}H_{42}$, (b) $C_{100}H_{202}$, and (c) $C_{1400}H_{2802}$ system, at four sliding velocities (from top to bottom) 0.3, 3, 10, and 100 m/s. Increasing the velocity results in a reduced effective viscosity; i.e., the thin films exhibit shear thinning.

The velocity dependence of the molecular dynamics data in Fig. 1 can be very well fitted by the formula $\eta_{\text{eff}} = B\dot{\gamma}^{-n}$ or

$$\log \eta_{\text{eff}} = C - n \log \dot{\gamma}, \quad (2)$$

where $C = \log B$. This is illustrated in Fig. 2 for two cases. The rest of the cases also show that a linear fit is appropriate with the exceptions of the results at very high temperatures. The $C_{100}H_{202}$ system shows the highest effective viscosity due to entanglement. The $C_{20}H_{42}$ is nearly

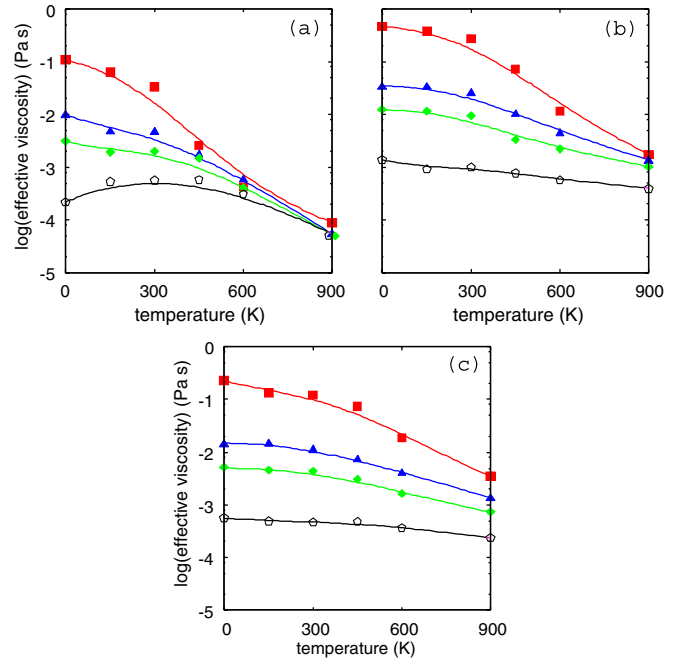


FIG. 1 (color online). The logarithm of the effective viscosity as a function of the temperature for (a) $C_{20}H_{42}$, (b) $C_{100}H_{202}$, and (c) $C_{1400}H_{2802}$ system at four sliding velocities: 0.3 m/s (■), 3 m/s (▲), 10 m/s (◆), and 100 m/s (◇).

liquid above 300 K and shows less entanglement whereas the sliding in $C_{1400}H_{2802}$ takes place at nearly one interface (see Fig. 6). We find that the parameters C and n depend on the temperature. The data points in Fig. 3 show the temperature dependence of the index n , while the solid lines are fits to the data points using the inverse sigmoidal curve:

$$n = \frac{1}{1 + (T/T_c)^\alpha}. \quad (3)$$

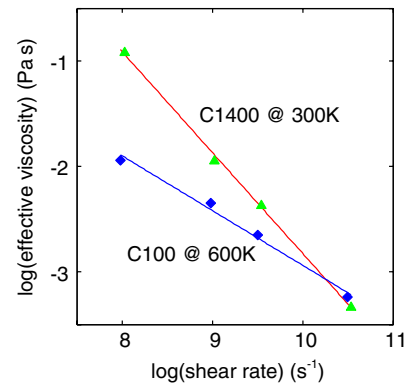


FIG. 2 (color online). The data points for the logarithm of the effective viscosity as a function of the logarithm of the shear rate at four sliding velocities: 0.3, 3, 10, and 100 m/s. The lines represent the linear fits to these data points. The systems are $C_{100}H_{202}$ at 600 K and $C_{1400}H_{2802}$ at 300 K. The values of the slope and the intercept of the $C_{100}H_{202}$ are -0.520 ± 0.038 and 2.260 ± 0.351 and for the $C_{1400}H_{2802}$ system -0.958 ± 0.024 and 6.753 ± 0.228 .

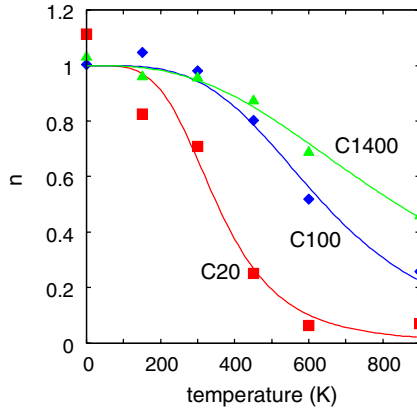


FIG. 3 (color online). Values of the parameter n in Eq. (2) as a function of the temperature for the systems $C_{20}H_{42}$, $C_{100}H_{202}$, and $C_{1400}H_{2802}$. The solid lines are fits to the numerical data using Eq. (3) with the parameters α and T_c given in Table I.

The parameters α and T_c are given in Table I. At low temperatures $n = 1$ as expected for dry friction. That is, the frictional shear stress $\sigma_f = \eta_{\text{eff}} \dot{\gamma} = 10^C \dot{\gamma}^{1-n}$ is independent of the shear rate when $n = 1$ as expected for dry friction at low temperatures (no thermally activated creep). At high temperature n approaches 0 as expected for a Newtonian fluid where the frictional shear stress is proportional to the shear rate.

The parameter C depends on the units used for η_{eff} and $\dot{\gamma}$ and here we assume SI units. Remarkably, we find a linear relation between C and n for all the systems and temperatures we have studied, see Fig. 4. The extrapolation of C to $n = 0$ gives $C \approx -3.8 \pm 0.2$. This gives the effective viscosity $\eta_{\text{eff}} = 10^C = (1.6\text{--}2.5) \times 10^{-4}$ Pa s. Experiments have shown [22] that for a wide range of fluids the fluid viscosity at the boiling point is $\approx 2.2 \times 10^{-4}$ Pa s, i.e., nearly the same as we deduce for confined fluids when extrapolating our data to $n = 0$ (Newtonian fluid). This result is extremely interesting but perhaps not entirely unexpected, as $n = 0$ corresponds to high temperatures where the separation between the solid walls (see Fig. 5 in which the sliding velocity is 10 m/s) is much larger than for lower temperatures and where the mobility of the fluid molecules may be similar to that in the bulk fluid close to the boiling point. The $C_{100}H_{202}$ film has a smaller film thickness than the other systems at low temperatures due to the fact that it only has 6 monolayers at these conditions. The other two systems have 7 monolayers at the same conditions and have thus thicker films. This matter has been discussed in Ref. [14].

TABLE I. Table showing the T_c and α in Eq. (3) for each system. The standard deviation of the fitting parameters is also indicated.

	$C_{20}H_{42}$	$C_{100}H_{202}$	$C_{1400}H_{2802}$
T_c (K)	353 ± 28	642 ± 22	840 ± 30
α (-)	4.09 ± 1.24	3.68 ± 0.46	2.79 ± 0.30

Figure 3 shows that the longer the hydrocarbon chain length, the higher the temperature necessary for the film to behave as a Newtonian fluid. This is also illustrated in Fig. 6 which shows the relative number of atoms traveling at velocities below v , as a function of v , for all three polymer films at 600 K. It can be seen that the velocity gradient is going from nearly Newtonian in the case of $C_{20}H_{42}$ to solidlike in the $C_{1400}H_{2802}$ case.

Note that for the hydrocarbons $C_{100}H_{202}$ and $C_{1400}H_{2802}$ at room temperature $n \approx 0.9$, as also found experimentally for many confined fluids. When $n \approx 0.9$, from Fig. 4 we get $C \approx 6$ which is a little larger than found experimentally (at room temperature) $C_{\text{exp}} \approx 5$. This may reflect a somewhat different confinement condition, e.g., differences in contact pressures (10 MPa in our study as compared to a few MPa in most of the experimental studies). However, in both cases the contact pressure is so small that one does not expect any significant dependency of the shear stress on the contact pressure [13], except if there is a pressure induced change in the number of confined layers.

It is well known that the viscosity of fluids at high pressures may be many orders of magnitude larger than at low pressures. Using the theory of activated processes, and assuming that a local molecular rearrangement in a fluid results in a local volume expansion, one expects an exponential dependence of viscosity η on the hydrostatic pressure p , $\eta = \eta_0 \exp(p/p_0)$, where typically (for hydrocarbons or polymer fluids) $p_0 \approx 10^8$ Pa (see, e.g., Refs. [23,24]). Here we are interested in (wetting) fluids confined between the surfaces of elastically soft solids, e.g., rubber. In this case, the pressure at the interface is usually at most of the order of the Young's modulus, which (for rubber) is less than 10^7 Pa. Thus, in most cases involving elastically soft materials, the viscosity can be considered as independent of the local pressure.

One of us has recently studied rubber friction on rough surfaces [25]. For unfilled styrene butadiene rubber we found the transfer of a thin smear film to the substrate. In

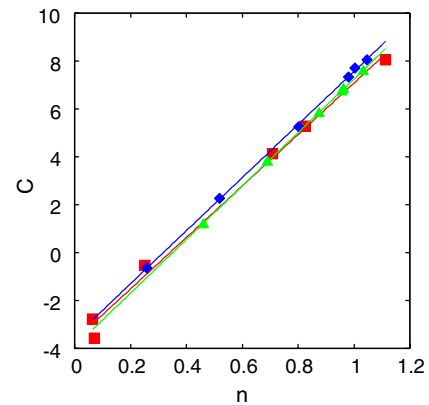


FIG. 4 (color online). The C parameter in Eq. (2) also follows a sigmoidal curve when the temperature is varied. This figure shows the dependence of C on n . All three systems $C_{20}H_{42}$, $C_{100}H_{202}$, and $C_{1400}H_{2802}$ are represented.

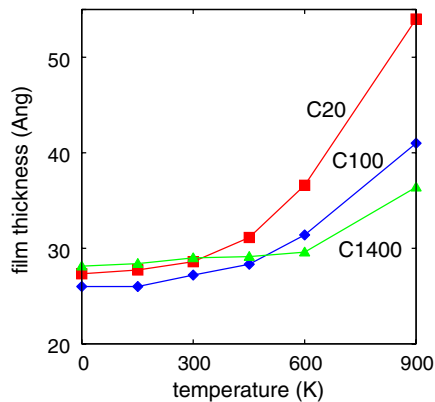


FIG. 5 (color online). The polymer film thickness as a function of the temperature. The sliding speed is 10 m/s. All three systems $C_{20}H_{42}$, $C_{100}H_{202}$, and $C_{1400}H_{2802}$ are represented.

this case, the shear during sliding may be localized to a thin (a few nanometer) interfacial layer which may exhibit frictional properties very similar to what we have observed in our simulations. Indeed, the experimental data indicated a frictional shear stress of the form predicted above with $n \approx 0.91$ and $10^C \approx 1.3 \times 10^5$ (in SI units), in close agreement with the result of our simulations.

To summarize, we have presented results of molecular dynamics calculations of friction performed for a block sliding on a substrate separated by ≈ 3 nm thick polymer films where the alkanes had 20, 100, and 1400 carbon atoms. In all cases, we found that the logarithm of the effective viscosity is proportional to the logarithm of the shear rate, $\log \eta_{\text{eff}} \approx C - n \log \dot{\gamma}$. The index n varies from 1 (solidlike friction) at very low temperatures to 0 (Newtonian liquid) at very high temperatures, following an inverse sigmoidal curve. The C parameter is proportional to n and as $n \rightarrow 0$, 10^C extrapolates to the viscosity of the bulk fluid at the boiling point. At room temperature, the parameters n and C have been found to be close to what has been observed experimentally for a large number of fluids.

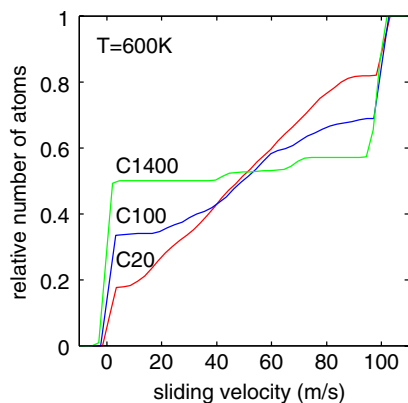


FIG. 6 (color online). The relative number of atoms moving at the velocity below v , as a function of v , at 600 K. It can be seen that the velocity gradient is going from nearly Newtonian in the case of $C_{20}H_{42}$ to solidlike in the $C_{1400}H_{2802}$ case.

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- [1] B. N. J. Persson and M. Scaraggi, *Eur. Phys. J. E* **34**, 113 (2011).
- [2] E. Kumacheva and J. Klein, *J. Chem. Phys.* **108**, 7010 (1998).
- [3] B. N. J. Persson and F. Mugele, *J. Phys. Condens. Matter* **16**, R295 (2004).
- [4] L. Bureau, *Phys. Rev. Lett.* **104**, 218302 (2010).
- [5] M. D. Demetriou *et al.*, *Phys. Rev. Lett.* **97**, 065502 (2006).
- [6] A. Furukawa, K. Kim, S. Saito, and H. Tanaka, *Phys. Rev. Lett.* **102**, 016001 (2009).
- [7] S. Yamada, *Tribol. Lett.* **13**, 167 (2002).
- [8] S. Granick, *Science* **253**, 1374 (1991).
- [9] G. Luengo, F.-J. Schmitt, R. Hill, and J. Israelachvili, *Macromolecules* **30**, 2482 (1997).
- [10] P. A. Thompson, G. S. Grest, and M. O. Robbins, *Phys. Rev. Lett.* **68**, 3448 (1992).
- [11] M. O. Robbins and A. R. C. Baljon, in *Microstructure and Microbiology of Polymer Surfaces* (American Chemical Society, Washington, DC, 2000), chap. 6.
- [12] Y. K. Yew, M. Minn, S. K. Sinha, and V. B. C. Tan, *Langmuir* **27**, 5891 (2011).
- [13] I. M. Sivebaek, V. N. Samoilov, and B. N. J. Persson, *Eur. Phys. J. E* **27**, 37 (2008).
- [14] I. M. Sivebaek, V. N. Samoilov, and B. N. J. Persson, *Langmuir* **26**, 8721 (2010).
- [15] B. N. J. Persson, V. N. Samoilov, S. Zilberman, and A. Nitzan, *J. Chem. Phys.* **117**, 3897 (2002).
- [16] I. M. Sivebaek, V. N. Samoilov, and B. N. J. Persson, *J. Chem. Phys.* **119**, 2314 (2003).
- [17] V. N. Samoilov and B. N. J. Persson, *J. Chem. Phys.* **120**, 1997 (2004).
- [18] W. L. Jorgensen, J. D. Madura, and C. J. Swenson, *J. Am. Chem. Soc.* **106**, 6638 (1984).
- [19] D. K. Dysthe, A. H. Fuchs, and B. Rousseau, *J. Chem. Phys.* **112**, 7581 (2000).
- [20] M. G. Martin and J. I. Siepmann, *J. Phys. Chem. B* **102**, 2569 (1998).
- [21] The temperature was obtained from the total kinetic energy but even at the highest sliding velocity 100 m/s the contribution from the center of mass motion to the temperature is not very large, corresponding to an increase in the temperature by only ~ 15 K, and much smaller at the lower sliding velocities (the contribution to the temperature from the translational motion scales as the square of the sliding velocity).
- [22] A. W. Francis, *Ind. Eng. Chem.* **49**, 1779 (1957).
- [23] H. Eyring, *J. Chem. Phys.* **4**, 283 (1936).
- [24] B. N. J. Persson, *Sliding Friction: Physical Principles and Applications* (Springer, Heidelberg, 2000), 2nd ed.
- [25] B. Lorenz, B. N. J. Persson, S. Dieluweit, and T. Tada, *Eur. Phys. J. E* **34**, 129 (2011).